

## Product datasheet for RC200310L3V

### OriGene Technologies, Inc.

9620 Medical Center Drive, Ste 200 Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com CN: techsupport@origene.cn

# STUB1 (NM\_005861) Human Tagged ORF Clone Lentiviral Particle

#### **Product data:**

**Product Type:** Lentiviral Particles

**Product Name:** STUB1 (NM\_005861) Human Tagged ORF Clone Lentiviral Particle

Symbol: STUB1

Synonyms: CHIP; HSPABP2; NY-CO-7; SCA48; SCAR16; SDCCAG7; UBOX1

Mammalian Cell

Selection:

Puromycin

**Vector:** pLenti-C-Myc-DDK-P2A-Puro (PS100092)

Tag: Myc-DDK
ACCN: NM 005861

ORF Size: 909 bp

**ORF Nucleotide** 

The ORF insert of this clone is exactly the same as(RC200310).

OTI Disclaimer:

Sequence:

The molecular sequence of this clone aligns with the gene accession number as a point of reference only. However, individual transcript sequences of the same gene can differ through naturally occurring variations (e.g. polymorphisms), each with its own valid existence. This clone is substantially in agreement with the reference, but a complete review of all prevailing

variants is recommended prior to use. More info

**OTI Annotation:** This clone was engineered to express the complete ORF with an expression tag. Expression

varies depending on the nature of the gene.

**RefSeq:** <u>NM 005861.2</u>

RefSeq Size: 1650 bp

RefSeq ORF: 912 bp

Locus ID: 10273

UniProt ID: Q9UNE7

Cytogenetics: 16p13.3

Domains: TPR, U-box

**Protein Families:** Druggable Genome





### STUB1 (NM\_005861) Human Tagged ORF Clone Lentiviral Particle - RC200310L3V

**Protein Pathways:** Ubiquitin mediated proteolysis

**MW:** 34.9 kDa

**Gene Summary:** This gene encodes a protein containing tetratricopeptide repeat and a U-box that functions

as a ubiquitin ligase/cochaperone. The encoded protein binds to and ubiquitinates shock cognate 71 kDa protein (Hspa8) and DNA polymerase beta (Polb), among other targets. Mutations in this gene cause spinocerebellar ataxia, autosomal recessive 16. Alternative splicing results in multiple transcript variants. There is a pseudogene for this gene on

chromosome 2. [provided by RefSeq, Jun 2014]